In the claims:

Following is a complete listing of the claims pending in the application, as amended:

- 1. (Currently amended) A topical delivery system, comprising:
- a gemini surfactant in admixture with a biologically active agent, wherein the delivery system, when in contact with skin or a mucosal membrane, provides a localized or systemic therapeutic effect.
- 2. (Currently amended) The delivery system according to claim 1, wherein the gemini surfactant is selected from the group consisting of an anionic gemini surfactant, a gemini cationic surfactant, a neutral gemini surfactant, an amphoteric gemini surfactant, ander mixtures thereof.
- 3. (Currently amended) The delivery system according to claim 1-or claim 2, wherein the gemini cationic surfactant has a hydrophobic tail comprising a C₃-C₃₀ alkyl group.
- 4. (Currently amended) The delivery system according to any one of claims 1-3claim 1, wherein the biologically active agent is a plasmid DNA.
- 5. (Previously presented) The delivery system according to claim 4, wherein the plasmid DNA comprises a gene encoding for interferon-γ.
- 6. (Currently amended) The delivery system according to any one of claims 1-5claim 1, wherein the delivery system includes further comprises one or more pharmaceutically-acceptable vehicles.
- 7. (Currently amended) The delivery system according to any one of claims 1-5claim 1, wherein the delivery system includes further comprises one or more supplements suitable for application for skin or mucosa.

- 8. (Currently amended) The delivery system according to claim 6-or 7, wherein the delivery system is <u>formulated to have a form selected from the group consisting in the form</u> of a cream, <u>a</u> lotion, <u>a</u> paste, <u>an</u> ointment, <u>a</u> foam, <u>a</u> gel, <u>a</u> lipid formulation, <u>an</u> emulsion, <u>a</u> solution, <u>orand a</u> suspension.
- 9. (Currently amended) The delivery system according to claim 8, further comprising one or more supplements selected from a neutral carrier erand a permeation enhancer.
- 10. (Previously presented) The delivery system according to claim 9, wherein the neutral carrier is selected from 1,2-dioleyl-sn-glycero-phosphatidylethanolamine (DOPE) or cholesterol.
- 11. (Previously presented) The delivery system according to claim 8, further comprising a compound selected from diethylene glycol monoethyl ether, polyglyceryl 3-diisostearate, PEG-8 caprylic and capric glycerides, and octyldodecyl myristate.
- 12.-19. (Cancelled)
- 20. (New) A method for treatment of a skin disorder, comprising topically delivering a delivery system according to claim 1.
- 21. (New) The method according to claim 20, wherein said delivering comprises delivering to a subject having a skin disorder selected from the group consisting of scleroderma, atopic dermatitis, and psoriasis.
- 22. (New) The method according to claim 20, wherein said delivering comprises delivering to a subject having a skin disorder of genetic origin.
- 23. (New) A method for treatment of a metabolic disease, comprising topically delivering a delivery system according to claim 1, wherein said metabolic disease is selected from the group consisting of gyrate atrophy, maternal hyperphenylalaninemia, familial hypercholesterolemia, and phenylketonuria.